

K896544 MD2 PERSONAL INFUSION SYSTEMMay 2, 1990
166 days to decisionK896544 · Product code: **FRN** · General Hospital
Source: <https://www.510kdatabase.net/k896544/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Nov 17, 1989
Decision date	May 2, 1990
Days to decision	166 days
Third-party review	No

APPLICANT

Company	Norfolk Medical Products, Inc.
Location	Walker, MI, US
Contact	MICHAEL J DALTON
510(k) history	20 submissions · 20 cleared · 1983-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k896544/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 18, 2026